Exhibit H

Drugs@FDA: FDA-Approved Drugs

f SHARE (HTTPS://WWW.FACEBOOK.COM/SHARER/SHARER.PHP?U=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/DAF/INDEX.CFM? EVENT=OVERVIEW.PROCESS&APPLNO=205029)

<u>▼ TWEET (HTTPS://TWITTER.COM/INTENT/TWEET/?TEXT=DRUGS@FDA: FDA-APPROVED</u>
DRUGS&URL=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/DAF/INDEX.CFM?EVENT=OVERVIEW.PROCESS&APPLNO=205029)

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■ EMAIL (MAILTO:?SUBJECT=DRUGS@FDA: FDA-APPROVED DRUGS&BODY=HTTPS://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/DAF/INDEX.CFM?EVENT=OVERVIEW.PROCESS&APPLNO=205029)

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New Drug Application (NDA): 205029

Company: BELCHER

■ EMAIL (MAILTO:?SUBJECT=DRUGS@FDA: FDA APPROVED DRUG
PRODUCTS&BODY=HTTP://WWW.ACCESSDATA.FDA.GOV/SCRIPTS/CDER/DAF/INDEX.CFM?
EVENT=OVERVIEW.PROCESS%26VARAPPLNO=205029)

Products on NDA 205029

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CSV Excel Print								
Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	TE Code	RLD		
EPINEPHRINE	EPINEPHRINE	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	SOLUTION; INTRAVENOUS, INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS	Prescription	None	Yes		

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Approval Date(s) and History, Letters, Labels, Reviews for NDA 205029

Original Approvals or Tentative Approvals

CSV E	excel Print				
Action Date	Submission	Action Type	Submission Classification	Review Priority; Orphan Status	Letters, Reviews, Labels, Patic
07/29/2014	07/29/2014 ORIG-1		Type 7 - Drug Already Marketed without Approved NDA	STANDARD	Label (PDF) (https://www.accessda Letter (PDF) (https://www.accessda Review (https://www.accessdata.fd

Showing 1 to 1 of 1 entries

Supplements

CSV	Ex	cel	Print			
Action Date		Submission		Supplement Categories or Approval Type	Letters, Reviews, Labels, Patient Package Insert	
05/18/20	2016 SUPPL-4				Labeling- Package Insert	Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/label/2016/2056 Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/appletter/2016/
02/11/20	02/11/2016 SUPPL-2		Efficacy-New Indication	Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/label/2016/2056 Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/appletter/2016/		
12/03/20	12/03/2015 SUPPL-3		Manufacturing (CMC)			
10/23/2015 SUPPL-1		Efficacy-New Indication	Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/2056 Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/appletter/2015/			

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Labels for NDA 205029

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